



Food and Drug Administration
10903 New Hampshire Avenue
Document Control Center - WO66-G609
Silver Spring, MD 20993-0002

May 31, 2016

AngioScore, Inc.
c/o Kimberley Kline
Regulatory Affairs Manager
5055 Brandin Court
Fremont, CA 94538

Re: K100303

Trade/Device Name: AngioSculpt PTA Scoring Catheter
Regulation Number: 21 CFR 870.1250
Regulation Name: Percutaneous catheter
Regulatory Class: Class II (two)
Product Code: PNO
Dated: February 2, 2010
Received: February 3, 2010

Dear Ms. Kline:

This letter corrects our substantially equivalent letter of March 22, 2010.

We have reviewed your Section 510(k) premarket notification of intent to market the device referenced above and have determined the device is substantially equivalent to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act) that do not require approval of a premarket approval application (PMA). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration. Please note: CDRH does not evaluate information related to contract liability warranties. We remind you, however, that device labeling must be truthful and not misleading.

If your device is classified (see above) into either class II (Special Controls) or class III (PMA), it may be subject to additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 898. In addition, FDA may publish further announcements concerning your device in the Federal Register.

Please be advised that FDA's issuance of a substantial equivalence determination does not mean that FDA has made a determination that your device complies with other requirements of the Act or any Federal statutes and regulations administered by other Federal agencies. You must

comply with all the Act's requirements, including, but not limited to: registration and listing (21 CFR Part 807); labeling (21 CFR Part 801); medical device reporting (reporting of medical device-related adverse events) (21 CFR 803); good manufacturing practice requirements as set forth in the quality systems (QS) regulation (21 CFR Part 820); and if applicable, the electronic product radiation control provisions (Sections 531-542 of the Act); 21 CFR 1000-1050.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801), please contact the Division of Industry and Consumer Education at its toll-free number (800) 638-2041 or (301) 796-7100 or at its Internet address

<http://www.fda.gov/MedicalDevices/ResourcesforYou/Industry/default.htm>. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21 CFR Part 807.97). For questions regarding the reporting of adverse events under the MDR regulation (21 CFR Part 803), please go to <http://www.fda.gov/MedicalDevices/Safety/ReportaProblem/default.htm> for the CDRH's Office of Surveillance and Biometrics/Division of Postmarket Surveillance.

You may obtain other general information on your responsibilities under the Act from the Division of Industry and Consumer Education at its toll-free number (800) 638-2041 or (301) 796-7100 or at its Internet address

<http://www.fda.gov/MedicalDevices/ResourcesforYou/Industry/default.htm>.

Sincerely yours,

Misti L. Malone -S

for Bram D. Zuckerman, M.D.

Director

Division of Cardiovascular Devices

Office of Device Evaluation

Center for Devices and Radiological Health

Enclosure

AngioScore, Inc
AngioSculpt® PTA Scoring Balloon Catheter

510(k) Premarket Notification

INDICATIONS FOR USE STATEMENT

510(k) Number (if known): K100303

Device Name: AngioSculpt® PTA Scoring Balloon Catheter

Indications for Use:

The AngioSculpt PTA Scoring Balloon Catheter is intended for dilatation of lesions in the iliac, femoral, ilio-femoral, popliteal, infra popliteal, and renal arteries, and for the treatment of obstructive lesions of native or synthetic arteriovenous dialysis fistulae. Not for use in the coronary or neuro-vasculature.

Prescription Use X
(Part 21 CFR 801 Subpart D)

AND/OR

Over-The-Counter Use _____
(21 CFR 801 Subpart C)

(PLEASE DO NOT WRITE BELOW THIS LINE-CONTINUE ON ANOTHER PAGE OF NEEDED)

Concurrence of CDRH, Office of Device Evaluation (ODE)

Diana R. V. Anne
(Division Sign-Off)
Division of Cardiovascular Devices

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510(k) Number K100303



AngioScore

510(k) Summary for the AngioSculpt Scoring Balloon Catheter

1. Submitter's Name / Contact Person

MAR 22 2010

Submitter: AngioScore, Inc.
5055 Brandin Court
Fremont, CA 94538

Contact Person: Kimberley Kline
Regulatory Affairs Manager
Phone: 510.933.7989
Fax: 510.933.7994

Summary Preparation Date: March 19, 2010

2. General Information

Trade Name: AngioSculpt® PTA Scoring Balloon Catheter

Common / Usual Name: Angioplasty catheter

Classification Name: Catheter, angioplasty, peripheral, transluminal

Product Codes: DQY and LIT

Predicate Devices: Cordis Aviator® PTA Dilatation Catheter (K013581)
Cordis Savvy® PTA Dilatation Catheter (K971010)
VascuTrak™ PTA Dilatation Catheter (K063657)
VascuTrak™ PTA Dilatation Catheter (K082343)
AngioSculpt® Scoring Balloon Catheter (K050629,
K072225, K080151, K081220, K082059, K091966)

3. Intended Use / Indications

The AngioSculpt PTA Scoring Balloon Catheter is intended for dilatation of lesions in the iliac, femoral, ilio-femoral, popliteal, infra popliteal, and renal arteries, and for the treatment of obstructive lesions of native or synthetic arteriovenous dialysis fistulae. Not for use in the coronary or neuro-vasculature.

4. Device Description

The AngioSculpt catheter is a standard two-lumen catheter with a scoring balloon near the distal tip. The distal end of the catheter has a conventional nylon-blend balloon with a nitinol scoring element that wraps around the balloon. The scoring element creates focal concentrations of dilating force which minimizes balloon slippage and assists with luminal expansion of stenotic arteries. The balloon has radiopaque markers to aid in positioning the balloon in the stenosis, and is designed to provide an expandable segment of known diameter and length at a specific pressure.

5. Substantial Equivalence Comparison

The AngioSculpt catheter intended use and that of the predicate devices, the Cordis Aviator PTA Dilatation Catheter, the Cordis Savvy PTA Dilatation Catheter, and the VascuTrak PTA Dilatation Catheter are substantially the same.

The indications for use of the subject device include the addition of renal arteries for the 510(k) cleared AngioSculpt PTA Scoring Balloon Catheter.

The AngioSculpt catheter shares the same fundamental design, scientific technology (operating principle), functional performance, and materials as the currently marketed AngioSculpt catheter. The subject device has similar design, operating principle, and functional performance as the Cordis Aviator and Savvy Dilatation Catheters, as well as the VascuTrak Dilatation Catheter.

Design verification and validation testing demonstrated adequate device performance and confirmed that no new questions of safety or effectiveness for peripheral balloon angioplasty devices were raised.

The inclusion of renal arteries in the indications for use does not raise new clinical risks. The anatomic and histologic features of renal arteries are substantially similar to those in the periphery, such as the infra-inguinal arteries. Both locations are similar in size and have similar fibro-fatty and calcified plaque formation.

The AngioSculpt PTA Scoring Balloon Catheter is therefore, substantially equivalent to the predicate devices.